

PATENT COOPERATION TREATY

WV

BUR

From the
INTERNATIONAL SEARCHING AUTHORITY

Roche Diagnostics GmbH
Patent Department Penzberg

PCT

To:

see form PCT/ISA/220

ASK	29. JUNI 2005		WN
BK			WJ
BUR	HH	HIL	MI SR

WRITTEN OPINION OF THE
INTERNATIONAL SEARCHING AUTHORITY
(PCT Rule 43bis.1)

Date of mailing
(day/month/year) see form PCT/ISA/210 (second sheet)

Applicant's or agent's file reference

see form PCT/ISA/220

FOR FURTHER ACTION

See paragraph 2 below

International application No.
PCT/EP2004/012459

International filing date (day/month/year)
04.11.2004

Priority date (day/month/year)
04.11.2003

International Patent Classification (IPC) or both national classification and IPC
G01N33/574, C12Q1/68

Applicant
ROCHE DIAGNOSTICS GMBH

Termin

04.09.05
(24.07.05) not. ✓

1. This opinion contains indications relating to the following items:

- ☒ Box No. I Basis of the opinion
- ☐ Box No. II Priority
- ☒ Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- ☒ Box No. IV Lack of unity of invention
- ☒ Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- ☐ Box No. VI Certain documents cited
- ☐ Box No. VII Certain defects in the international application
- ☒ Box No. VIII Certain observations on the international application

2. FURTHER ACTION

If a demand for international preliminary examination is made, this opinion will usually be considered to be a written opinion of the International Preliminary Examining Authority ("IPEA"). However, this does not apply where the applicant chooses an Authority other than this one to be the IPEA and the chosen IPEA has notified the International Bureau under Rule 66.1bis(b) that written opinions of this International Searching Authority will not be so considered.

If this opinion is, as provided above, considered to be a written opinion of the IPEA, the applicant is invited to submit to the IPEA a written reply together, where appropriate, with amendments, before the expiration of three months from the date of mailing of Form PCT/ISA/220 or before the expiration of 22 months from the priority date, whichever expires later.

For further options, see Form PCT/ISA/220.

3. For further details, see notes to Form PCT/ISA/220.

Name and mailing address of the ISA:



European Patent Office
D-80298 Munich
Tel. +49 89 2399 - 0 Tx: 523656 epmu d
Fax: +49 89 2399 - 4465

Authorized Officer

Thumb, W

Telephone No. +49 89 2399-7350



Best Available Copy

WRITTEN OPINION OF THE
INTERNATIONAL SEARCHING AUTHORITYInternational application No.
PCT/EP2004/012459

AP20 Rec'd PCT/PTO 14 APR 2006

Box No. I Basis of the opinion

1. With regard to the **language**, this opinion has been established on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.
☐ This opinion has been established on the basis of a translation from the original language into the following language , which is the language of a translation furnished for the purposes of international search. (under Rules 12.3 and 23.1(b)).
2. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application and necessary to the claimed invention, this opinion has been established on the basis of:
 - a. type of material:
☐ a sequence listing
☐ table(s) related to the sequence listing
 - b. format of material:
☐ in written format
☐ in computer readable form
 - c. time of filing/furnishing:
☐ contained in the international application as filed.
☐ filed together with the international application in computer readable form.
☐ furnished subsequently to this Authority for the purposes of search.
3. ☐ In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished.
4. Additional comments:

Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non obvious), or to be industrially applicable have not been examined in respect of:

- ☐ the entire international application,
- ☒ claims Nos. 1-27 (partially)

because:

- ☒ the said international application, or the said claims Nos. 22-27 relate to the following subject matter which does not require an international preliminary examination (*specify*):

see separate sheet

- ☐ the description, claims or drawings (*indicate particular elements below*) or said claims Nos. are so unclear that no meaningful opinion could be formed (*specify*):
- ☐ the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.
- ☒ no international search report has been established for the whole application or for said claims Nos. 1-27 (partially)
- ☐ the nucleotide and/or amino acid sequence listing does not comply with the standard provided for in Annex C of the Administrative Instructions in that:
 - the written form ☐ has not been furnished
 - ☐ does not comply with the standard
 - the computer readable form ☐ has not been furnished
 - ☐ does not comply with the standard
- ☐ the tables related to the nucleotide and/or amino acid sequence listing, if in computer readable form only, do not comply with the technical requirements provided for in Annex C-bis of the Administrative Instructions.
- ☐ See separate sheet for further details

**WRITTEN OPINION OF THE
INTERNATIONAL SEARCHING AUTHORITY**

International application No.
PCT/EP2004/012459

Box No. IV Lack of unity of invention

1. ☒ In response to the invitation (Form PCT/ISA/206) to pay additional fees, the applicant has:
- ☐ paid additional fees.
 - ☐ paid additional fees under protest.
 - ☒ not paid additional fees.
2. ☐ This Authority found that the requirement of unity of invention is not complied with and chose not to invite the applicant to pay additional fees.
3. This Authority considers that the requirement of unity of invention in accordance with Rule 13.1, 13.2 and 13.3 is
- ☐ complied with
 - ☒ not complied with for the following reasons:
see separate sheet
4. Consequently, this report has been established in respect of the following parts of the international application:
- ☐ all parts.
 - ☒ the parts relating to claims Nos. 1-27 (partially).

Box No. V Reasoned statement under Rule 43b/s.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes: Claims	1-18, 20, 21
	No: Claims	19
Inventive step (IS)	Yes: Claims	
	No: Claims	1-21
Industrial applicability (IA)	Yes: Claims	1-21
	No: Claims	

2. Citations and explanations

see separate sheet

Box No. VIII Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

see separate sheet

**WRITTEN OPINION OF THE
INTERNATIONAL SEARCHING
AUTHORITY (SEPARATE SHEET)**

PCT/EP2004/012459

AP20 Rec'd PCT/PTO 14 APR 2006

Re Item III

Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

Claims 26 and 27 pertain to a reference data bank for distinguishing immunologically defined ALL subtypes Pro-B-ALL, c-ALL, Pre-B-ALL, c-ALL/Pre-B-ALL, mature B-ALL, precursor B-ALL, Pro-T-ALL, Pre-T-ALL, cortical T-ALL, mature T-ALL and/or T-ALL in a sample.

A data bank as such is characterised only by data contained in said data bank, which are considered to be a mere presentation of information. No international preliminary examination is carried out for the subject-matter of said claims pursuing the provisions of Rule 67.1(v) PCT.

It should further be noted that the technical information presented under points (a) and (b) of claim 26 is related to the method of constructing said data bank and is therefore no characterising technical feature of the data bank as such, claimed in claim 26.

An analogous argumentation also applies to the subject-matter of claims 22-25. Presentation of information is not patentable whether the claims are directed to the presentation of the information per se or to apparatus for presenting the information which are solely defined by the information recorded (see also the Preliminary Examination Guidelines, Chapter 9, Item 9.12). Again, the method for obtaining a data bank does not define the data bank as such.

Re Item IV

Lack of unity of invention

1. The application lacks unity within the meaning of Rule 13.1 PCT.
The problem to be solved in the present application is the provision of markers for distinguishing immunologically defined ALL subtypes Pro-B-ALL, c-ALL, Pre-B-ALL, c-ALL/Pre-B-ALL, mature B-ALL, precursor B-ALL, Pro-T-ALL, Pre-T-ALL, cortical T-ALL, mature T-ALL and/or T-ALL in a sample.
The single general concept which can be identified a priori as linking the various inventions and which forms a solution to the above problem relates to the use of "markers for ALL subtypes". The use of marker genes/nucleotides disclosed in tables

1 and 2 form 1050 different solutions to the above problem.

However, the concept of using marker genes for distinguishing different ALL subtypes is known in the art.

Kohlmann et al., Genes, Chromosomes & Cancer (2003) Vol. 37, pp. 396-405 (**D1**) discloses a method for distinguishing precursor-B-ALL (subtype t(8;14)) from precursor T-ALL (see page 402, col. 1, paragraph 2) using expression of ADA as a marker (table 3, page 404). This marker is disclosed for the same purpose in the present application in table 2.8, position 38.

A subtype with MLL translocation is distinguished from the remainder using expression of VLDLR as a marker (table 3, page 404). This marker is disclosed for the same purpose in table 1.6, position 22 of the present application (definition of the subtypes in Example 1, page 23 of the present application).

Document Tsutsumi et al., Cancer Research (1993) Vol. 53, pp. 4882-4887 (**D2**) discloses markers for distinguishing ALL with MLL gene arrangements (i.e. Pro-B-ALL according to the definition given in Example 1, page 23 of the present application) from other ALL subtypes, including CD44 and MEIS1 (see the abstract and page 4885, col. 2, lines 7-10). It is noted that MLL-Re-ALL can be diagnosed from gene expression profiles...

Said markers are included for the same purpose in table 1.6, positions 18 and 10, respectively.

Yeoh et al., Cancer Cell (2002) Vol. 1, pp. 133-143 (**D3**) also discloses MEIS1 as a specific marker for MLL-ALL (i.e. Pro-B-ALL) (page 136, col. 2, paragraph 1), as in table 1.6 of the present application (see preceding paragraph). It is also noted in D3 that gene expression profiles can identify each of the prognostically important leukemia subtypes (see abstract). Affymetrix microarrays are used for the expression analysis (see Experimental procedures, page 142).

Document WO-A-03/039443 (**D4**) discloses several markers for distinguishing ALL subtypes, identified using gene expression analysis (page 71 and 72).

Document WO-A-03/083140 (**D5**) pertains to characterisation of ALL subtypes using

differential gene expression analysis. In table 66, D5 discloses several sets of markers for distinguishing ALL subtypes including subtypes with MLL translocations (i.e. Pro-B-ALL according to page 23, Example 1 of the present application). Said markers include MEIS1, CD72, C20orf103, and CD44, which are all included in table 1.6 of the present application.

In the light of D1-D5, each document taken alone, the above identified single general concept is not novel and inventive and thus cannot be the single general inventive concept as required by Rule 13.1 PCT.

The present invention is thus considered not to fulfil the requirements of unity as laid down in Rule 13.1 PCT.

No other technical features could be identified that form a technical relationship among each of the separate inventions claimed and which could be considered as same or corresponding special technical features within the meaning of Rule 13.2 PCT.

The first invention was searched, namely methods relating to distinguishing immunologically defined ALL subtypes, in particular mature B-ALL from other subtypes using CD99 as a marker; kits and apparatus for distinguishing mature B-ALL from other subtypes using said marker.

2. The Examining Authority considers that the following separate inventions or groups of inventions are not so linked as to form a single general inventive concept:

Invention 1: Claims 1-27 (all partially)

A method for distinguishing mature B-ALL from all other subtypes of immunologically defined ALL subtypes, the method comprising determining the expression level of the marker CD99. Use of said marker for the manufacture of

a diagnostic. A diagnostic kit containing said marker and an apparatus comprising a reference data bank, wherein the reference data bank is obtainable by determining the expression level of CD99.

Inventions 2-1050: Claims 1-27 (all partially)

Methods for distinguishing immunologically defined ALL subtypes Pro-B-ALL, c-ALL, Pre-B-ALL, c-ALL/Pre-B-ALL, mature B-ALL, precursor B-ALL, Pro-T-ALL, Pre-T-ALL, cortical T-ALL, mature T-ALL and/or T-ALL and methods for distinguishing specific subtypes against all other subtypes and against each other, the method comprising determining individually the expression level of the markers listed in tables 1.1, positions 2-50, tables 1.2-1.6 and in table 2. Use of said markers for the manufacture of diagnostics. Diagnostic kits containing said markers and apparatus comprising a reference data bank; wherein the reference data bank is obtainable by determining the expression levels of said markers.

The following assessment of novelty and inventive step will only pertain to subject-matter for which a search report has been established, i.e. invention 1.

Re Item V

Reasoned statement with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

- 1.1 Claim 19 does not meet the requirements of Article 33(2) PCT.
Claim 19 refers to kit for distinguishing leukemia subtypes containing at least CD99. This must be construed as meaning merely a reagent suitable for carrying out the method. The intended use of a product is not a technical feature of the product per se. Therefore, commercially available microarrays, such as the U133 microarrays of Affymetrix, comprising CD99 specific probes are novelty-destroying for the subject-matter of claim 19 within the meaning of Article 33(2) PCT.

- 1.2 Claims 1-18 and 20-21 are novel within the meaning of Article 33(2) PCT, since the prior art does not teach the use of CD99 as a marker for distinguishing mature B-ALL from all other subtypes of immunologically defined ALL subtypes, or kits and apparatus comprising a reference for immunologically defined ALL subtypes based on CD99 expression.
- 2.1 Claim 1 does not meet the requirements of Article 33(3) PCT.
Documents D1-D5, each of which could be considered to represent the most relevant state of the art, disclose markers for distinguishing immunologically defined ALL subtypes.
The underlying objective technical problem may therefore be seen in providing a further marker for distinguishing ALL subtypes.
As already pointed out under item IV, 1. above, the use of differential gene expression analysis using microarrays of gene probes for defining ALL subtypes is described in detail in documents D1-D5. In addition, several other documents pertain to the concept of identifying gene expression profiles in order to characterise ALL subtypes (see for example Chen et al., Blood (2001) Vol. 97(7), pp. 2115-2120 (D6): abstract; Kohlmann et al. (2002) Blood, Vol. 100(11), Abstract No. 1212 (D7): the whole document).
Moreover, methods for classifying samples based on gene expression data have become common general knowledge in the art, also in the field of leukemia diagnosis (see for example EP-A-1 043 676 (D8), the whole document; Kohlmann et al. (2002) Blood, Vol. 100(11); Abstract No. 4287 (D9); Deutsch, J.M. (2003) Bioinformatics, Vol.19(1), pp.45-52 (D10)).
The above referred-to documents represent a non-exhaustive list of documents dealing with the identification of marker genes indicative of a specific leukemia subtype.
In addition, CD99 (a synonym for MIC2) has already been described as a marker for ALL subtypes in document D5, table 34, position 24, and in document D7. These documents also describes differentiation of ALL subtypes using analysis of differential gene expression on Affymetrix microarrays.

In particular documents D1-D5 contain direct pointers that it is possible to identify gene markers which are specific for a certain ALL subtype and thus enable an

unambiguous identification of said subtypes.

Moreover, the use of CD99 as a marker does not appear to be associated with an unexpected and surprising technical effect in view of the above-cited documents which could confer an inventive step compared to other markers identified by gene expression profiling using standard microarray technology.

It would therefore be obvious for a person skilled in the art to use differential gene expression based on microarray analysis in order to identify further markers, e.g. CD99, for specific ALL subtypes in view of state of the art as exemplified in documents D1-D10 in order to solve the above-stated problem.

Hence, claim 1 cannot be considered as being inventive within the meaning of Article 33(3) PCT.

- 2.2 ~~Claims 2-21 refer to standard embodiments in the art of microarray analysis and diagnostics and do not add technical features which would confer an inventive activity.~~

Claims 2-21 thus do also not meet the requirements of Article 33(3) PCT.

3. Should the objection under Rule 67.1(v) be overcome, the applicant is referred to documents Dugas et al. (2002), In silico biology, Vol. 2, pp. 383-391 (**D11**) and Dugas et al. (2001) Leukemia, Vol. 15, pp. 1805-1810 (**D12**), which disclose databases containing data from patients suffering from leukemia. Said data include characterisation of subtypes, and correlation of cytogenetic findings with, e.g., microarray data (D12: page 1807, col. 2; **D11**: the whole document). Therefore, claims pertaining to the generation of reference databases for the analysis of leukemia subtypes based on gene expression data could not be considered as being novel (Article 33(2) PCT).

Re Item VIII

Certain observations on the international application

1. In order to avoid any unclarity within the meaning of Article 6 PCT, abbreviations should be defined the first time they are mentioned in the claims.
2. Notwithstanding the objection of lack of unity raised under item 2. above, claim 1 does not meet the requirements of Article 6 PCT. The excessive use of "and/or" for defining various possible embodiments in claim 1 as well as the introduction of an enormous number of possible marker combinations through the use of the term "at least one polynucleotide" in each of said possible embodiments, the claim lacks conciseness, contrary to the requirements of Article 6 PCT.

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ BLACK BORDERS
- ☐ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES
- ☒ FADED TEXT OR DRAWING
- ☒ BLURRED OR ILLEGIBLE TEXT OR DRAWING
- ☐ SKEWED/SLANTED IMAGES
- ☐ COLOR OR BLACK AND WHITE PHOTOGRAPHS
- ☐ GRAY SCALE DOCUMENTS
- ☒ LINES OR MARKS ON ORIGINAL DOCUMENT
- ☒ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY
- ☐ OTHER: _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.